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STIC Search Report

STIC Database Tracking Number: 93878

TO: James Pasterczyk Location: CP3 8A11

May 13, 2003

Case Serial Number: 09359359

From: Kathleen Fuller Location: EIC 1700

CP3/4 3D62

Phone: 308-4290

Kathleen.Fuller@uspto.gov

Search Notes

I believe I found the exact compounds in the claims. I did not limit them by utility but printed all 6 references.

There is a mistake in the preparation for example 273 in the attached claims you gave me. One of the ingredients should be a benzoyl or benzoic acid compound.



Access DB# 93878

SEARCH REQUEST FORM

Scientific and Technical Information Center

Requester's Full Name: Art Unit: 1755	Dhans William Pysteria	Examiner #: 71209 Date: 5/12/03
Mail Box and Bldg/Roor	n Location: CA3- X A1	PAPER DISK E-MA Results Format Preferred (circle): PAPER DISK E-MA
If more than one search	n is submitted, please p	prioritize searches in order of need.
Please provide a detailed state	ment of the search tomic and	******************
Include the elected species or	structures, keywords, synonyn	describe as specifically as possible the subject matter to be searched. ns, acronyms, and registry numbers, and combine with the concept or
known. Please attach a copy o	e any terms that may have a sp	us, acronyms, and registry numbers, and combine with the concept or becall meaning. Give examples or relevant citations, authors, etc., if
17 -	and sover sheet, pertinent class	ms, and abstract.
Title of Invention:		
Inventors (please provide ful	l names):	
Earliest Priority Filing Da	te:	
*For Sequence Searches Only *	Please include all and a control	
appropriate serial number.	reuse include all pertinent infori	mation (parent, child, divisional, or issued patent numbers) along with the
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AFF USE ONLY	******	***************
rcher: K. Fuller	Type of Search	Vendors and cost where applicable
cher Phone #:	NA Sequence (#)	
rcher Location:	AA Sequence (#)	
Searcher Picked Up:	Structure (#)	Questel/Orbit
Completed:	Bibliographic	Dr.Link
cher Prep & Review Time: 20	Litigation	Lexis/Nexis
	Fulltext	Sequence Systems
cal Prep Time:	Date of the	
ne Time: 7	Patent Family	WWW/Internet

, EIC1700

Search Results Feedback Form (Optional)



The search results generated for your recent request are attached. If you have any questions or comments (compliments or complaints) about the scope or the results of the search, please contact the EIC searcher who conducted the search or contact:

Kathleen Fuller, Team Leader, 308-4290, CP3/4 3D62

Volu	ntary Results Feedback Form
>	I am an examiner in Workgroup: Example: [1713]
>	Relevant prior art found, search results used as follows:
	102 rejection
	103 rejection
	Cited as being of interest.
	Helped examiner better understand the invention.
	Helped examiner better understand the state of the art in their technology.
	Types of relevant prior art found:
	Foreign Patent(s)
	Non-Patent Literature (journal articles, conference proceedings, new product announcements etc.)
>	Relevant prior art not found:
	Results verified the lack of relevant prior art (helped determine patentability).
	Search results were not useful in determining patentability or understanding the invention
)ther	Comments:

=> FILE REG

1

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> D QUE

L28

150 SEA FILE=REGISTRY ABB=ON (19481-82-4/BI OR 213453-19-1/BI OR 9003-21-8/BI OR 9003-53-6/BI OR 10025-73-7/BI OR 10031-26-2/BI OR 10049-05-5/BI OR 102-82-9/BI OR 104-81-4/BI OR 106-95-6/BI OR 106173-87-9/BI OR 106826-12-4/BI OR 106911-77-7/BI OR 107227-34-9/BI OR 108150-11-4/BI OR 108501-18-4/BI OR 108501-19-5/BI OR 108548-96-5/BI OR 110-18-9/BI OR 110772-34-4/BI OR 110807-37-9/BI OR 111-40-0/BI OR 1116-76-3/BI OR 111740-42-2/BI OR 112965-31-8/BI OR 121264-61-7/BI OR 121876-18-4/BI OR 123-72-8/BI OR 124-63-0/BI OR 1313-13-9/BI OR 148-24-3/BI OR

7

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1643-19-2/BI OR 166664-50-2/BI OR 17201-43-3/BI OR 17639-93-9/B
                I OR 17678-99-8/BI OR 18301-66-1/BI OR 188065-74-9/BI OR
                197142-57-7/BI OR 197142-58-8/BI OR 197142-59-9/BI OR 197251-85
                -7/BI OR 204580-80-3/BI OR 2052-01-9/BI OR 205379-28-8/BI OR
                20769-85-1/BI OR 208446-93-9/BI OR 2085-33-8/BI OR 212697-80-8/
                BI OR 213137-90-7/BI OR 213453-06-6/BI OR 213453-12-4/BI OR
                213453-13-5/BI OR 213453-14-6/BI OR 213453-21-5/BI OR 213555-59
                -0/BI OR 2212-32-0/BI OR 222734-87-4/BI OR 23426-63-3/BI OR
                24457-21-4/BI OR 25014-41-9/BI OR 25038-76-0/BI OR 25038-78-2/B
                I OR 25067-63-4/BI OR 25084-99-5/BI OR 25154-86-3/BI OR
                25232-27-3/BI OR 25249-16-5/BI OR 25266-62-0/BI OR 2549-51-1/BI
                 OR 25767-47-9/BI OR 26022-14-0/BI OR 26374-91-4/BI OR
                26490-65-3/BI OR 26588-80-7/BI OR 29158-71-2/BI OR 2916-14-5/BI
                OR 29263-94-3/BI OR 3012-37-1/BI OR 3030-47-5/BI OR 30323-87-6
                /BI OR 30811-69-9/BI OR 3083-10-1/BI OR 33527-91-2/BI OR
                34946-82-2/BI OR 366-18-7/BI OR 37234-97-2/BI OR 39149-80-9/BI
                OR 40704-75-4/BI OR 4097-89-6/BI OR 41203-22-9/BI OR 4328-13-6/
                BI OR 49864-98-4/BI OR 503445-26-9/BI OR 503445-27-0/BI OR
                5061-21-2/BI OR 51359-78-5/BI OR 535-11-5/BI OR 5445-17-0/BI
                OR 5468-93-9/BI OR 56467-21-1/BI OR 56905-18-1/BI OR 585-71-7/B
                I OR 589-15-1/BI OR 590-17-0/BI
L35
              3 SEA FILE=REGISTRY ABB=ON L28 AND AZO
L36
              1 SEA FILE=REGISTRY ABB=ON L35 AND 2/NR
L37
             4 SEA FILE=HCAPLUS ABB=ON L36
L38
             1 SEA FILE=REGISTRY ABB=ON L28 AND VINYL ACETATE
L42
          13190 SEA FILE=REGISTRY ABB=ON 108-05-4/CRN
L43
             1 SEA FILE=REGISTRY ABB=ON 108-05-4
L44
             1 SEA FILE=REGISTRY ABB=ON 371771-79-8
L45
             1 SEA FILE=REGISTRY ABB=ON 371771-77-6/CRN
L46
              1 SEA FILE=REGISTRY ABB=ON 213453-06-6
L50
          88095 SEA FILE=HCAPLUS ABB=ON L38 OR L42 OR L43
L51
             4 SEA FILE=HCAPLUS ABB=ON L37 AND L50
L53
             1 SEA FILE=REGISTRY ABB=ON 213453-03-3
L54
             1 SEA FILE=REGISTRY ABB=ON
                                         213453-02-2/CRN
L55
             1 SEA FILE=REGISTRY ABB=ON 213453-02-2
L56
              4 SEA FILE=HCAPLUS ABB=ON (L53 OR L54 OR L55) OR (L44 OR L45 OR
                L46)
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=> D L57 ALL 1-6 HITSTR

- L57 ANSWER 1 OF 6 HCAPLUS COPYRIGHT 2003 ACS
- AN 2003:255121 HCAPLUS
- DN 138:272092

L57

TI Atom or group transfer radical polymerization in the presence of transition metals applicante

6 SEA FILE=HCAPLUS ABB=ON L51 OR L56

- IN Matyjaszewski, Krzysztof; Gaynor, Scott G.; Coca, Simion
- PA Carnegie Mellon University, USA
- SO U.S., 90 pp., Cont.-in-part of U.S. 6,407,187. CODEN: USXXAM
- DTPatent
- LΑ English
- IC ICM C08F004-06 ICS C08F004-40; C08F004-42
- 526090000; 526113000; 526118000; 526135000; 526172000; 526328000; NCL 526335000; 526346000; 526347000
- CC 35-3 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 67

FAN.CNT 6 PATENT NO.		KIND	DATE	AP	PLICATION NO.	DATE	
PI	US	6541580	B1	20030401	US	1999-369157	19990806
	US	5763548	Α	19980609	US	1995-414415	19950331
	US	6538091	B1	20030325	US	1998-18554	19980204
	US	6407187	B1	20020618	US	1998-34187	19980303
	US	6512060	B1	20030128	US	1999-359591	19990723
	US	2002183473	A1	20021205	US	2001-34908	20011221
	US	2002193538	A1	20021219	US	2002-98052	20020313
PRAI	US	1995-414415	A3	19950331			
	US	1997-39543P	P	19970311			
	US	1997-41620P	P	19970402			
	US	1998-18554	A3	19980204			
	US	1998-34187	A2	19980303			
	US	1995-559309	A3	19951115			
	US	1999-369157	A2	19990806			
	US	2000-534827	A2	20000323			
	US	2000-257738P	P	20001222			
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AB A process for ATRP polymn. and coupling of mols. by radical processes is provided, wherein improvements are provided by using transition metal of zero oxidn. state in place of or in addn. to transition metal complexes to give improved control over mol. wt., mol. wt. distribution and compns. of the products formed. Alternatively, these improvements are achieved by using mixed transition metal compd. systems in which 1 of the transition metals is in a higher of 2 available oxidn. state and the other is in a lower of 2 available oxidn. states, wherein the 2 metals are different. Alternatively, these improvements are achieved by using compds. of Fe, Mn, Cr, or Cu that can participate in a reversible redox cycle with .gtoreq.1 of initiators, dormant polymer chain ends, and growing polymer chain ends. Thus, heating 10 mg Fe powder, 69 mg PPh3, 1 mL styrene, and 12 .mu.L 1-phenylethyl bromide 9 h at 110.degree. gave 70% polymer with Mn 6780 and Mw/Mn 1.19.

ST ATRP zero valent transition metal catalyst; polystyrene manuf iron triphenylphosphine phenylethyl bromide catalyst

IT Polysiloxanes, preparation

RL: IMF (Industrial manufacture); PREP (Preparation)
(atom or group transfer radical polymn. in presence of transition metal compds. that participate in reversible redox cycles with initiators, dormant polymer chain ends, or growing polymer chain ends)

IT Transition metals, uses

RL: CAT (Catalyst use); USES (Uses)

(atom or group transfer radical polymn. in presence of zero valent transition metals)

IT Transition metal complexes

RL: CAT (Catalyst use); USES (Uses)

(atom or group transfer radical polymn. in presence of zero valent transition metals and(or) transition metal compds.)

IT Polymerization catalysts

(atom transfer, radical; atom or group transfer radical polymn. in presence of zero valent transition metals and(or) transition metal compds.)

IT Boranes

Phosphines

RL: CAT (Catalyst use); USES (Uses)

(cocatalyst; atom or group transfer radical polymn. in presence of zero valent transition metals)

```
IT
    Dendritic polymers
    RL: IMF (Industrial manufacture); PREP (Preparation)
        (hyperbranched; atom or group transfer radical polymn. for manuf. of
       hyperbranched polymers from (bromopropionyloxy)ethyl acrylate)
ΙT
    Carboxylic acids, uses
    RL: CAT (Catalyst use); USES (Uses)
        (salts, cocatalyst; atom or group transfer radical polymn. in presence
       of zero valent transition metals)
IT
    598-54-9, Cuprous acetate
                                7447-39-4, Cupric chloride, uses
    Chromic chloride
    RL: CAT (Catalyst use); USES (Uses)
        (atom or group transfer radical polymn. in presence of mixed transition
       metal compd. system in which 1 of the transition metals is in higher of
       2 available oxidn. states)
                                      25249-16-5P, Poly-2-hydroxyethyl
IT
    25014-41-9P, Polyacrylonitrile
                   26588-80-7P, Butyl acrylate-2-hydroxyethyl
    methacrylate
    methacrylate-methyl methacrylate-styrene copolymer
    RL: IMF (Industrial manufacture); PREP (Preparation)
        (atom or group transfer radical polymn. in presence of mixed transition
       metal compd. system in which 1 of the transition metals is in higher of
       2 available oxidn. states)
ΙT
    1313-13-9, Manganese dioxide, uses
                                         7773-01-5, Manganese chloride
    10049-05-5, Chromium dichloride 26490-65-3, Cuprous hexafluorophosphate
    37234-97-2
                 68986-76-5, Cuprous 2-thiophenecarboxylate
    RL: CAT (Catalyst use); USES (Uses)
        (atom or group transfer radical polymn. in presence of transition metal
        compds. that participate in reversible redox cycles with initiators,
       dormant polymer chain ends, or growing polymer chain ends)
ΙT
    79-07-2DP, 2-Chloroacetamide, reaction products with polystyrene
    104-81-4DP, 4-Methylbenzyl bromide, reaction products with polystyrene
    106-95-6DP, Allyl bromide, reaction products with polymethyl acrylate
    589-15-1DP, 4-Bromobenzyl bromide, reaction products with polystyrene
    590-17-0DP, Bromoacetonitrile, reaction products with polystyrene
    2549-51-1DP, Vinyl chloroacetate, reaction products with polystyrene
    2916-14-5DP, Allyl chloroacetate, reaction products with polystyrene
    5061-21-2DP, reaction products with polystyrene 9003-21-8DP, Polymethyl
    acrylate, functionalized
                              9003-49-0P, Polybutyl acrylate
                                                               9003-54-7P,
    Acrylonitrile-styrene copolymer
                                     17201-43-3DP, 4-Cyanobenzyl bromide,
    reaction products with polystyrene
                                        19481-82-4DP, 2-Bromopropionitrile,
    reaction products with polystyrene <a>25067-63-4P</a>, Methyl
    acrylate-vinyl acetate copolymer 25154-86-3P, Poly[2-
     (dimethylamino)ethyl methacrylate]
                                         25232-27-3P, Poly-tert-butyl acrylate
    25266-62-0P, Polyallyl acrylate 25767-47-9P, Butyl acrylate-styrene
    copolymer
                26022-14-0P, Poly-2-hydroxyethyl acrylate 26374-91-4P,
    Polyglycidyl acrylate
                           29158-71-2P, Poly[2-(trimethylsiloxy)ethyl
                    30323-87-6P, Polyisobornyl acrylate
    methacrylate]
                                                          30811-69-9P,
                         39149-80-9DP, tert-Butyl 2-bromopropionate, reaction
    Polyvinyl acrylate
    products with polystyrene
                                40704-75-4P, Poly-N-(2-
    hydroxypropyl) methacrylamide
                                   56467-21-1P, Butyl acrylate-3-
     (trimethoxysily1)propyl methacrylate copolymer
                                                    86293-61-0P,
    N-Cyclohexylmaleimide-styrene copolymer
                                              106173-87-9P, 2-Hydroxyethyl
    methacrylate-styrene graft copolymer 106826-12-4P, Ethylene-styrene
    graft copolymer
                      106911-77-7P, Methyl methacrylate-styrene block
    copolymer
                107227-34-9P, Acrylonitrile-isobutylene alternating copolymer
    108150-11-4P, Methyl acrylate-methyl methacrylate-block copolymer
    108501-18-4P, Butyl acrylate-methyl methacrylate block copolymer
    108501-19-5P, Butyl acrylate-methyl methacrylate-styrene block copolymer
```

108548-96-5P, Hexamethylcyclotrisiloxane-styrene graft copolymer

TT

ΙŢ

IT

IT

110772-34-4P, Butyl acrylate-styrene block copolymer 110807-37-9P, Ethylene-methyl methacrylate graft copolymer 111740-42-2P, Methyl acrylate-styrene block copolymer 112965-31-8P, Acrylonitrile-butyl acrylate-styrene block copolymer 121264-61-7P, Butyl acrylate-2-hydroxyethyl acrylate block copolymer 121876-18-4P, Isobutylene-isoprene-styrene graft copolymer 166664-50-2P, Butyl acrylate-isobutylene alternating copolymer 197142-57-7P, Dicyclopentadiene-styrene block copolymer 197142-58-8P, Methyl 197142-59-9P, Dicyclopentadieneacrylate-norbornene block copolymer methyl acrylate block copolymer 197251-85-7P, Norbornene-styrene block 208446-93-9DP, 2-Hydroxyethyl 2-bromopropionate, reaction products with polystyrene 212697-80-8P, Poly[2-(trimethylsiloxy)ethyl 213453-12-4P, Styrene-vinyl chloride-vinyl chloroacetate graft acrylate] 213453-13-5P, Butyl acrylate-2-ethylhexyl acrylatecopolymer 3,3,4,4,5,5,6,6,7,7,8,8,9,9,10,10,10-heptadecafluorodecyl acrylate block copolymer 213453-14-6P, Isobornyl acrylate-isobutene-methyl acrylate-methyl methacrylate-styrene block copolymer 213555-59-0P, Acrylonitrile-methyl acrylate block copolymer 222734-87-4P, Isobutene-p-methylstyrene-styrene graft copolymer 503445-26-9P, Isobutene-isobornyl acrylate-p-methylstyrene graft copolymer 503445-27-0P, Isobornyl acrylate-isobutene-isoprene graft copolymer RL: IMF (Industrial manufacture); PREP (Preparation) (atom or group transfer radical polymn. in presence of transition metal compds. that participate in reversible redox cycles with initiators, dormant polymer chain ends, or growing polymer chain ends) 9003-53-6DP, Polystyrene, reaction products with glycidyl 2-bromopropionate 85673-60-5DP, reaction products with polystyrene RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent) (atom or group transfer radical polymn. in presence of transition metal compds. that participate in reversible redox cycles with initiators, dormant polymer chain ends, or growing polymer chain ends) 7439-88-5, Iridium, uses 7439-89-6, Iron, uses 7439-96-5, Manganese, 7440-02-0, Nickel, uses 7440-05-3, Palladium, uses 7440-06-4. 7440-15-5, Rhenium, uses 7440-16-6, Rhodium, uses Platinum, uses 7440-19-9, Samarium, uses 7440-22-4, 7440-18-8, Ruthenium, uses Silver, uses 7440-47-3, Chromium, uses 7440-50-8, Copper, uses 7440-66-6, Zinc, uses RL: CAT (Catalyst use); USES (Uses) (atom or group transfer radical polymn. in presence of zero valent transition metals) 9003-53-6P, Polystyrene 9003-21-8P, Polymethyl acrylate 9011-14-7P, 188065-74-9P, Poly[2-(2-bromopropionyloxy)ethyl acrylate] 213453-06-6P, Poly[2-(2-bromopropionyloxy)ethyl methacrylate] RL: IMF (Industrial manufacture); PREP (Preparation) (atom or group transfer radical polymn. in presence of zero valent transition metals) 49864-98-4P, Hexakis[4-(hydroxymethyl)phenoxy]cyclotriphosphazene RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent) (cocatalyst precursor; atom or group transfer radical polymn. in presence of transition metal compds. that participate in reversible redox cycles with initiators, dormant polymer chain ends, or growing. polymer chain ends) 111-40-0, Diethylenetriamine 64-18-6, Formic acid, reactions 123-72-8, Butyraldehyde 4097-89-6, Tren RL: RCT (Reactant); RACT (Reactant or reagent) (cocatalyst precursor; atom or group transfer radical polymn. in

```
presence of transition metal compds. that participate in reversible
        redox cycles with initiators, dormant polymer chain ends, or growing
        polymer chain ends)
     2085-33-8, Aluminum 8-Hydroxyquinolinate
ΙT
     RL: CAT (Catalyst use); USES (Uses)
        (cocatalyst; atom or group transfer radical polymn. in presence of
        mixed transition metal compd. system in which 1 of the transition
        metals is in higher of 2 available oxidn. states)
IT
     80-58-0, 2-Bromobutyric acid
                                    94-36-0, Benzoyl peroxide, uses
    N,N,N',N'-Tetramethylethylenediamine 124-63-0, Methanesulfonyl chloride
                                          535-11-5, Ethyl 2-bromopropionate
     148-24-3, 8-Hydroxyguinoline, uses
                                       998-40-3, Tributylphosphine
     672-65-1, 1-Phenylethyl chloride
     1116-76-3, Trioctylamine
                                1643-19-2, Tetrabutylammonium bromide
     2052-01-9, 2-Bromoisobutyric acid
                                         2212-32-0, 2-[[2-
     (Dimethylamino)ethyl]methylamino]ethanol
                                                3012-37-1, Benzyl thiocyanate
                         4328-13-6, Tetrahexylammonium bromide
                                                                 17639-93-9,
     3030-47-5, PMDETA
     Methyl 2-chloropropionate
                                 18301-66-1, Trimethylsilyl 2-bromobutyrate
                                              41203-22-9, 1,4,8,11-Tetramethyl-
     24457-21-4, tert-Butyl 2-bromobutyrate
     1,4,8,11-tetraazacyclotetradecane
                                        56905-18-1, Methyl 2-iodopropionate
     72914-19-3, 4,4'-Di-tert-butyl-2,2'-bipyridine
                                                     82280-42-0,
     Hexakis [4-(bromomethyl)phenoxy]cyclotriphosphazene
                                                           213137-90-7,
     tert-Butyldimethylsilyl 2-bromobutyrate
     RL: CAT (Catalyst use); USES (Uses)
        (cocatalyst; atom or group transfer radical polymn. in presence of
        transition metal compds. that participate in reversible redox cycles
        with initiators, dormant polymer chain ends, or growing polymer chain
        ends)
     5468-93-9P, 1,2-Bis(2-bromopropionyloxy)ethane
                                                       17678-99-8P
IT
     33527-91-2P, Tris[2-(dimethylamino)ethyl]amine
                                                      204580-80-3P
     RL: CAT (Catalyst use); IMF (Industrial manufacture); PREP (Preparation);
     USES (Uses)
        (cocatalyst; atom or group transfer radical polymn. in presence of
        transition metal compds. that participate in reversible redox cycles
        with initiators, dormant polymer chain ends, or growing polymer chain
        ends)
     68-12-2, DMF, uses
                          98-59-9, p-Toluenesulfonyl chloride
IT
     Tributylamine 366-18-7, 2,2'-Bipyridine 585-71-7, 1-Phenylethyl
               600-00-0, Ethyl 2-bromoisobutyrate
                                                    603-35-0,
                               3083-10-1, 1,1,4,7,10,10-
     Triphenylphosphine, uses
     Hexamethyltriethylenetetramine 5445-17-0, Methyl 2-bromopropionate
     7787-70-4, Cuprous bromide 7789-45-9, Cupric bromide Ferrous bromide 10031-26-2, Ferric bromide 19481-82
                                                    19481-82-4,
     .alpha.-Bromopropionitrile
                                 23426-63-3, Methyl 2-bromoisobutyrate
     29263-94-3, Diethyl methylbromomalonate
                                              34946-82-2, Copper ditriflate
     72230-93-4, 4,4'-Bis(5-nonyl)-2,2'-bipyridine
     RL: CAT (Catalyst use); USES (Uses)
        (cocatalyst; atom or group transfer radical polymn. in presence of zero
        valent transition metals)
ΙT
     213453-21-5P
     RL: CAT (Catalyst use); IMF (Industrial manufacture); PREP (Preparation);
     USES (Uses)
        (initiator; atom or group transfer radical polymn. in presence of
        transition metal compds. that participate in reversible redox cycles
        with initiators, dormant polymer chain ends, or growing polymer chain
ΙT
     9016-00-6DP, Hexamethylcyclotrisiloxane homopolymer, sru, derivs.
```

25084-99-5DP, Hexamethylcyclotrisiloxane homopolymer, derivs.

RL: IMF (Industrial manufacture); PREP (Preparation)

(macroinitiator or macromonomers; atom or group transfer radical polymn. in presence of transition metal compds. that participate in reversible redox cycles with initiators, dormant polymer chain ends, or growing polymer chain ends)

IT 61551-69-7P

RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)

(macroinitiator precursor; atom or group transfer radical polymn. in presence of transition metal compds. that participate in reversible redox cycles with initiators, dormant polymer chain ends, or growing polymer chain ends)

IT 213453-19-1

RL: RCT (Reactant); RACT (Reactant or reagent)
(macroinitiator precursor; atom or group transfer radical polymn. in
presence of transition metal compds. that participate in reversible
redox cycles with initiators, dormant polymer chain ends, or growing
polymer chain ends)

IT 20769-85-1DP, 2-Bromoisobutyryl bromide, reaction products with polyhydroxyethyl methacrylate 81601-52-7DP, reaction products with hexamethylcyclotrisiloxane homopolymer RL: CAT (Catalyst use); IMF (Industrial manufacture); PREP (Preparation); USES (Uses)

(macroinitiator; atom or group transfer radical polymn. in presence of transition metal compds. that participate in reversible redox cycles with initiators, dormant polymer chain ends, or growing polymer chain

IT 213453-19-1DP, reaction products with polystyrene
RL: IMF (Industrial manufacture); PREP (Preparation)
 (macroinitiator; atom or group transfer radical polymn. in presence of transition metal compds. that participate in reversible redox cycles with initiators, dormant polymer chain ends, or growing polymer chain

9011-12-5DP, Isobutene-styrene copolymer, chlorinated 25038-76-0DP, Polynorbornene, reaction products with bromomethylbenzaldehyde 25038-78-2DP, Polydicyclopentadiene, reaction products with bromomethylbenzaldehyde 51359-78-5DP, 4-Bromomethylbenzaldehyde, reaction products with polynorbornene

RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)

(macroinitiator; atom or group transfer radical polymn. in presence of transition metal compds. that participate in reversible redox cycles with initiators, dormant polymer chain ends, or growing polymer chain ends)

IT 205379-28-8DP, reaction products with hexamethylcyclotrisiloxane homopolymer

RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)

(macromonomer; atom or group transfer radical polymn. in presence of transition metal compds. that participate in reversible redox cycles with initiators, dormant polymer chain ends, or growing polymer chain ends)

IT 9003-69-4P, Polydivinylbenzene

RL: IMF (Industrial manufacture); PREP (Preparation)
(star; atom or group transfer radical polymn. in presence of transition metal compds. that participate in reversible redox cycles with initiators, dormant polymer chain ends, or growing polymer chain ends)

IT 7631-86-9, Silica, uses

RL: CAT (Catalyst use); USES (Uses)

(support; atom or group transfer radical polymn. in presence of transition metal compds. that participate in reversible redox cycles with initiators, dormant polymer chain ends, or growing polymer chain ends)

RE.CNT 45 THERE ARE 45 CITED REFERENCES AVAILABLE FOR THIS RECORD RE

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- IT 25067-63-4P, Methyl acrylate-vinyl acetate copolymer
 - RL: IMF (Industrial manufacture); PREP (Preparation)

(atom or group transfer radical polymn. in presence of transition metal compds. that participate in reversible redox cycles with initiators, dormant polymer chain ends, or growing polymer chain ends)

RN 25067-63-4 HCAPLUS

```
PASTERCZYK 09/359359
                         Page 9
    2-Propenoic acid, methyl ester, polymer with ethenyl acetate (9CI) (CA
CN
    INDEX NAME)
    CM
         1
    CRN
        108-05-4
    CMF C4 H6 O2
Aco-CH-CH2
    CM
         2
    CRN 96-33-3
    CMF C4 H6 O2
    0
MeO-C-CH-CH2
    213453-06-6P, Poly[2-(2-bromopropionyloxy)ethyl methacrylate]
ΙT
    RL: IMF (Industrial manufacture); PREP (Preparation)
        (atom or group transfer radical polymn. in presence of zero valent
        transition metals)
    213453-06-6 HCAPLUS
RN
CN
    2-Propenoic acid, 2-methyl-, 2-(2-bromo-1-oxopropoxy)ethyl ester,
    homopolymer (9CI) (CA INDEX NAME)
    CM
         1
    CRN 213453-05-5
    CMF C9 H13 Br O4
 H<sub>2</sub>C
Me-C-C-O-CH_2-CH_2-O-C-CH-Me
IT
    213<u>453-21-</u>5P
    RL: CAT (Catalyst use); IMF (Industrial manufacture); PREP (Preparation);
    USES (Uses)
        (initiator; atom or group transfer radical polymn. in presence of
        transition metal compds. that participate in reversible redox cycles
       with initiators, dormant polymer chain ends, or growing polymer chain
        ends)
RN
    213453-21-5 HCAPLUS
CN
    Benzoic acid, 4-(chloromethyl)-, azobis[(2,2-dimethyl-1-oxo-2,1-
    ethanediyl)imino-2,1-ethanediyl] ester (9CI) (CA INDEX NAME)
          AMCBP spample 273
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KATHLEEN FULLER EIC 1700/PARKER LAW 308-4290

AMCBP elected. species

PAGE 1-B

CH₂Cl

ANSWER 2 OF 6 HCAPLUS COPYRIGHT 2003 ACS L57

AN 2001:629056 HCAPLUS

DN 135:344821

TΙ Synthesis and characterization of hyperbranched polyacrylates in the presence of a tetrafunctional initiator with higher reactivity than monomer by self-condensing vinyl polymerization

Hong, C.-Y.; Pan, C.-Y. ΑU

Department of Polymer Science and Engineering, University of Science and Technology of China, Hefei, Anhui, 230026, Peop. Rep. China Polyme((2001) 42(23), 9385-9391 CODEN: POLMAG: ISSN: 0032-3861

PB Elsevier Science Ltd.

DT Journal

LΑ English

CC 35-4 (Chemistry of Synthetic High Polymers)

AB A series of hyperbranched polyacrylates were synthesized by self-condensing vinyl polymn. (SCVP) of 2-[(2-bromobutyryl)oxy]ethyl acrylate (BBEA). A tetrafunctional atom transfer radical polymn. (ATRP) initiator (THABI) capped with bromoisobutyrate, which has higher reactivity than the initiating site of monomer, was used as core-forming mols. The structure and properties of the obtained polymers were characterized by NMR and SEC/RALLS/DV/RI. The effect of the tetrafunctional initiator on mol. wt. and mol. wt. distribution of polymers was studied.

ST atom transfer radical polymn initiator hyperbranched polyacrylate synthesis; self condensing vinyl polymn hyperbranched polyacrylate synthesis

TТ Polymerization

Polymerization catalysts

(atom transfer, radical; for synthesis of hyperbranched polyacrylates)

TΤ Dendritic polymers

> RL: SPN (Synthetic preparation); PREP (Preparation) (hyperbranched; synthesis of hyperbranched polyacrylates by self-condensing vinyl polymn.)

366-18-7, 2,2'-Bipyridine IT 11129-27-4, Copper bromide

RL: CAT (Catalyst use); USES (Uses) (initiation catalyst; in synthesis of hyperbranched polyacrylates by self-condensing vinyl polymn.)

```
371771-78-7P
IT
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
        (initiator; in synthesis of hyperbranched polyacrylates by
        self-condensing vinyl polymn.)
ΙT
     371771-77-6P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (monomer; in synthesis of hyperbranched polyacrylates in presence of a
        tetrafunctional initiator)
     20469-89-0, 2-Bromo-2-methylpropanoyl chloride
IT
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reactant for initiator; in synthesis of hyperbranched polyacrylates in
        presence of a tetrafunctional initiator)
IT
     182919-44-4P, 6,6-Bis (5-hydroxyl-2-oxapentyl)-4,8-dioxaundecane-1,11-diol
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (reactant for initiator; in synthesis of hyperbranched polyacrylates in
        presence of a tetrafunctional initiator)
                                   219901-16-3, 2-Hydroxyethyl 2-bromobutyrate
TΤ
     814-68-6, Acryloyl chloride
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reactant for monomer; in synthesis of hyperbranched polyacrylates in
        presence of a tetrafunctional initiator)
TΨ
     371771-79-8P
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (synthesis of hyperbranched polyacrylates by self-condensing vinyl
        polymn.)
RE.CNT
              THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD
        27
RE
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IT
     371771-79-8P
```

RL: SPN (Synthetic preparation); PREP (Preparation)

(synthesis of hyperbranched polyacrylates by self-condensing vinyl polymn.)

RN 371771-79-8 HCAPLUS

CN Butanoic acid, 2-bromo-, 2-[(1-oxo-2-propenyl)oxy]ethyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 371771-77-6 CMF C9 H13 Br O4 #282

elected.

L57 ANSWER 3 OF 6 HCAPLUS COPYRIGHT 2003 ACS

AN 1999:619836 HCAPLUS

DN 131:337428

TI Block copolymerizations of vinyl acetate by combination of conventional and atom transfer radical polymerization

AU Paik, Hyun-jong; Teodorescu, Mircea; Xia, Jianhui; Matyjaszewski, Krzysztof

CS Center for Macromolecular Engineering Department of Chemistry, Carnegie Mellon University, Pittsburgh, PA, 15213, USA

Macromolecules (1999), 32(21), 7023-7031

CODEN: MAMOBX; ISSN: 0024-9297

PB American Chemical Society

DT Journal

LA English

CC 35-4 (Chemistry of Synthetic High Polymers)

AB Four different methods of block copolymn., combining atom transfer radical polymn. (ATRP) and conventional radical polymn., were studied. The first two methods employed azo compds. contg. activated halogen atoms. 2,2'-Azobis[2-methyl-N-[2-[4-(chloromethyl)benzoyloxy]ethyl]propionamide] was used to initiate the polymn. of vinyl acetate (VAc) at 90 .degree.C. The resulting pVAc with a Cl terminal group (Mn = 47 900; Mw/Mn = 2.21) was subsequently used as a macroinitiator for styrene (St) to yield pVAc-b-pSt (Mn = 91 600; Mw/Mn = 1.80). In the second method, 2,2'-azobis[2-methyl-N-[2-(2-bromoisobutyryloxy)ethyl]propionamide] was first used to polymerize Bu acrylate (BA) at 30 .degree.C in the presence of CuBr/tris[2-(dimethylamino)ethyl]amine. The pBA (Mn = 7500; Mw/Mn = 1.15) with the preserved central azo unit was dissolved in VAc and extended to a block copolymer (Mn = 41 800; Mw/Mn = 3.56). Alternatively, ATRP has been combined with a redox initiated system. VAc was polymd. in the presence of CCl4/Fe(OAc)2/N,N,N',N'',N''-pentamethyldiethylenetriamine to yield pVAc with trichloromethyl end groups (Mn = 3600; Mw/Mn = 1.81). The polymer obtained was dissolved in styrene and block copolymd. by ATRP to form pVAc-b-pSt (Mn = 24 300; Mw/Mn = 1.42). In the last method, pBA with a bromine end group (Mn = 2460; Mw/Mn = 1.32) as prepd. by ATRP was dissolved in VAc together with CuBr/1,4,8,11-tetramethyl-1,4,8,11tetraazacyclotetradecane to initiate VAc polymn. A block copolymer with Mn = 4450 and Mw/Mn = 2.58 was prepd. In the presence of 20 mol % of CuBr2, the polydispersity was further reduced to 1.73.

ST vinyl acetate block radical polymn; atom transfer block polymn; styrene vinyl acetate block polymer; butyl acrylate vinyl acetate block copolymer

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IT
     Polymerization
        (block, group-transfer; block polymn. by combination of conventional
        and atom transfer radical methods)
     Polymerization
TΤ
        (block, radical; by combination of conventional and atom transfer
        radical methods)
TΤ
     Polymerization catalysts
        (block; in block polymn. by combination of conventional and atom
        transfer radical methods)
     Chain transfer agents
ΙT
        (in block polymn. by combination of conventional and atom transfer
        radical methods)
IT
     Macromonomers
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (in block polymn. by combination of conventional and atom transfer
        radical methods)
     Polymerization catalysts
TT
        (redox; in block polymn. by combination of conventional and atom
        transfer radical methods)
IT
     7758-89-6, Cuprous chloride
                                   7787-70-4, Cuprous bromide
                      33527-91-2, Tris[2-(dimethylamino)ethyl]amine
     Cupric bromide
     41203-22-9, 1,4,8,11-Tetramethyl-1,4,8,11-tetraazacyclotetradecane
     72230-93-4
     RL: CAT (Catalyst use); USES (Uses)
        (catalyst for block polymn. by combination of conventional and atom
        transfer radical methods)
     213453-19-1P, 2,2'-Azobis[2-methyl-N-[2-(2-bromoisobutyryloxy)ethyl]propio
IT
     namide] 213453-21-5P, 2,2'-Azobis[2-methyl-N-[2-[4-
     (chloromethyl)benzoyloxy]ethyl]propionamide]
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
        (catalyst for block polymn. by combination of conventional and atom
        transfer radical methods)
IT
     20769-85-1, 2-Bromoisobutyryl bromide 61551-69-7, 2,2'-Azobis[2-methyl-N-
     (2-hydroxyethyl)propionamide]
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (catalyst starting material; catalyst for block polymn. by combination
        of conventional and atom transfer radical methods)
     56-23-5, uses
RL: CAT (Catalyst use); USES (Uses)
IT
        (chain-transfer agent; catalysts in block polymn. by combination of
        conventional and atom transfer radical methods)
     107948-08-3P, Styrene-vinyl acetate block copolymer
IΤ
     110772-34-4P, Butyl acrylate-styrene block copolymer 135911-78-3P
     , Butyl acrylate-vinyl acetate block copolymer
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (prepn. by combination of conventional and atom transfer radical
        polymn.)
IΤ
     3030-47-5, PMDETA
                         3094-87-9, Ferrous acetate
     RL: CAT (Catalyst use); USES (Uses)
        (redox; catalyst for block polymn. by combination of conventional and
        atom transfer radical methods)
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- 213453-21-5P, 2,2'-Azobis[2-methyl-N-[2-[4-
 - (chloromethyl)benzoyloxy]ethyl]propionamide]
 - RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)
 - (catalyst for block polymn. by combination of conventional and atom transfer radical methods)
- RN 213453-21-5 HCAPLUS
- Benzoic acid, 4-(chloromethyl)-, azobis[(2,2-dimethyl-1-oxo-2,1-CNethanediyl)imino-2,1-ethanediyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 107948-08-3 HCAPLUS

CN Acetic acid ethenyl ester, polymer with ethenylbenzene, block (9CI) (CA INDEX NAME)

CM 1

CRN 108-05-4 CMF C4 H6 O2

AcO-CH=CH₂

CM 2

CRN 100-42-5 CMF C8 H8

 $H_2C == CH - Ph$

CM 1

CRN 141-32-2 CMF C7 H12 O2 О || n-BuO-C-CH== CH2

CM 2

CRN 108-05-4 CMF C4 H6 O2

 $AcO-CH=CH_2$

L57 ANSWER 4 OF 6 HCAPLUS COPYRIGHT 2003 ACS

AN 1999:558887 HCAPLUS

DN 132:166575

TI Use of difunctional azo initiators in the block copolymerization by combination of conventional and atom transfer radical polymerization

AU Paik, Hyun-jong; Matyjaszewski, Krzysztof

CS Center for Macromolecular Engineering Department of Chemistry, Carnegie Mellon University, Pittsburgh, PA, 15213, USA

SO Polymer Preprints (American Chemical Society, Division of Polymer Chemistry) (1999), 40(2), 436-437
CODEN: ACPPAY: TSSN: 0032-3934

PB American Chemical Society, Division of Polymer Chemistry

DT Journal

LA English

CC 35-3 (Chemistry of Synthetic High Polymers)

AB Block copolymers were prepd. using difunctional initiators having two functional groups. One of the functional groups is an azo group for initiation of conventional radical polymn. The other functional group is activated halogen for initiation of atom transfer radical polymn. (ATRP). Block copolymn. was performed using first either ATRP or conventional radical polymn., followed by the other method. Poly(Bu acrylate)-b-polystyrene, poly(vinyl acetate)-b-polystyrene and poly(Bu acrylate)-b-poly(vinyl acetate) were prepd. and characterized. The initiation efficiency of the (macro)azo initiator was studied.

ST azo initiator block radical polymn; catalyst polymn difunctional azo compd; atom transfer radical polymn azo initiator

IT Polymerization

(atom transfer, radical; use of difunctional azo initiators in block copolymn. by combination of conventional and atom transfer radical polymn.)

IT Polymerization

(radical; use of difunctional azo initiators in block copolymn. by combination of conventional and atom transfer radical polymn.)

IT Polymerization catalysts

(use of difunctional azo initiators in block copolymn. by combination of conventional and atom transfer radical polymn.)

IT 213453-19-1P, 2,2'-Azobis[2-methyl-N-(2-(2-bromoisobutyryloxy)ethyl)propio
namide 213453-21-5P, 2,2'-Azobis[2-methyl-N-(2-(4chloromethylbenzoyloxy)ethyl)propionamide
RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
USES (Uses)

(polymn. catalyst; use of difunctional azo initiators in block copolymn. by combination of conventional and atom transfer radical polymn.)

IT 20769-85-1, 2-Bromoisobutyryl bromide 61551-69-7, 2,2'-Azobis[2-methyl-N-(2-hydroxyethyl)propionamide

RL: RCT (Reactant); RACT (Reactant or reagent)

(reactant; in prepn. of difunctional azo initiators for block copolymn. by combination of conventional and atom transfer radical polymn.)

IT 107948-08-3P, Styrene-vinyl acetate block copolymer

110772-34-4P, Butyl acrylate-styrene block copolymer 135911-78-3P

, Butyl acrylate-vinyl acetate block copolymer

RL: SPN (Synthetic preparation); PREP (Preparation)

(use of difunctional azo initiators in block copolymn. by combination of conventional and atom transfer radical polymn.)

RE.CNT 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD RE

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- (3) Destarac, M; Polym Prepr (Am Chem Soc, Div Polym Chem) 1998, V39(2), P568 HCAPLUS
- (4) Kato, M; Macromolecules 1995, V28, P1721 HCAPLUS
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- (8) Matyjaszewski, K; J Am Chem Soc 1997, V119, P674 HCAPLUS
- (9) Moad, G; The Chemistry of Free-Radical Polymerization 1995
- (10) Percec, V; Macromolecules 1995, V28, P7970 HCAPLUS
- (11) Wang, J; J Am Chem Soc 1995, V117, P5614 HCAPLUS
- (12) Xia, J; Macromolecules 1998, V31, P5958 HCAPLUS

IT 213453-21-5p, 2,2'-Azobis[2-methyl-N-(2-(4chloromethylbenzoyloxy)ethyl)propionamide

RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);

USES (Uses)
 (polymn. catalyst; use of difunctional azo initiators in block
 copolymn. by combination of conventional and atom transfer radical
 polymn.)

RN 213453-21-5 HCAPLUS

CN Benzoic acid, 4-(chloromethyl)-, azobis[(2,2-dimethyl-1-oxo-2,1-ethanediyl)imino-2,1-ethanediyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

AMCBP

PAGE 1-B

107948-08-3P, Styrene-vinyl acetate block copolymer IT 135911-78-3P, Butyl acrylate-vinyl acetate block copolymer RL: SPN (Synthetic preparation); PREP (Preparation) (use of difunctional azo initiators in block copolymn. by combination of conventional and atom transfer radical polymn.) 107948-08-3 HCAPLUS RNAcetic acid ethenyl ester, polymer with ethenylbenzene, block (9CI) (CA CN INDEX NAME) CM 1 CRN 108-05-4 CMF C4 H6 O2 Aco-CH-CH2

CM 2

CRN 100-42-5 CMF C8 H8

 $H_2C = CH - Ph$

135911-78-3 HCAPLUS CN 2-Propenoic acid, butyl ester, polymer with ethenyl acetate, block (9CI) (CA INDEX NAME)

CM 1

CRN 141-32-2 C7 H12 O2 CMF

n-BuO-C-CH=CH2

CM 2

CRN 108-05-4 CMF C4 H6 O2

AcO-CH=CH2

L57 ANSWER 5 OF 6 HCAPLUS COPYRIGHT 2003 ACS AN 1998:807168 HCAPLUS 130:95895

DN

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Preparation of hyperbranched polyacrylates by atom-transfer radical
TΙ
     polymerization. Part 4. The use of zero-valent copper
     Matyjaszewski, Krzysztof; Pyun, Jeffrey; Gaynor, Scott G.
ΑU
ĊS
     Dep. Chemistry, Carnegie Mellon Univ., Pittsburgh, PA, 15213, USA
     Macromolecular Rapid Communications (1998), 19(12), 665-670
so
     CODEN: MRCOE3; ISSN: 1022-1336
PΒ
     Wiley-VCH Verlag GmbH
DT
     Journal
LΑ
     English
     35-3 (Chemistry of Synthetic High Polymers)
CC
     Section cross-reference(s): 36
AB
     The addn. of zero-valent Cu to the self-condensing vinyl polymn. (SCVP) of
     novel AB* (meth)acrylic monomers using atom-transfer radical polymn.
     (ATRP) catalyst systems has allowed for their successful polymn. Polymn.
     under homogeneous and heterogeneous catalyst conditions without addnl.
    Cu(0) were unsuccessful. The catalyst system that was designed comprised
     of Cu(I) bromide, 4,4'-bis(5-nonyl)-2,2'-bipyridine, and Cu(0) metal
     (powder or turning). From 1H NMR spectroscopy, the degree of branching was estd. for the acrylic polymers, DB = 0.48. The degree of branching
     could not be detd. for methacrylates by this method due to overlapping
     signals in the 1H NMR spectra.
ST
     hyperbranched polyacrylate radical polymn copper powder
ΙT
     Polymer chains
        (branching; prepn. of hyperbranched polyacrylates by atom-transfer
        radical polymn. by using zero-valent copper catalyst)
IT
     Dendritic polymers
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (prepn. of hyperbranched polyacrylates by atom-transfer radical polymn.
        by using zero-valent copper catalyst)
ΙT
     Polymerization catalysts
        (radical; prepn. of hyperbranched polyacrylates by atom-transfer
        radical polymn. by using zero-valent copper catalyst)
ΙT
     188065-74-9P 213453-03-3P 213453-06-6P
                                              213453-09-9P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (hyperbranched; prepn. of hyperbranched polyacrylates by atom-transfer
        radical polymn. by using zero-valent copper catalyst)
IT
     7440-50-8, Copper, uses 7758-89-6, Copper monochloride
                                                                  72230-93-4,
     4,4'-Bis(5-nonyl)-2,2'-bipyridine
     RL: CAT (Catalyst use); USES (Uses)
        (prepn. of hyperbranched polyacrylates by atom-transfer radical polymn.
        by using zero-valent copper catalyst)
RE.CNT
       17
              THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
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(2) Gaynor, S; ACS Symposium Series 1998, V685, P396 HCAPLUS
(3) Gaynor, S; Macromolecules 1996, V29, P1079 HCAPLUS
(4) Gaynor, S; Macromolecules 1997, V30, P5192
(5) Gaynor, S; Macromolecules 1997, V30, P7034
(6) Gaynor, S; Macromolecules 1997, V30, P7042
(7) Hawker, C; J Am Chem Soc 1995, V117, P10763 HCAPLUS
(8) Lu, P; Macromolecules 1996, V29, P8583 HCAPLUS
(9) Matyjaszewski, K; ACS Symposium Series 1998, V685, P258 HCAPLUS
(10) Matyjaszewski, K; Macromolecules 1995, V28, P7901
(11) Matyjaszewski, K; Macromolecules 1997, V30, P7348 HCAPLUS
(12) Matyjaszewski, K; Macromolecules 1998, V31, P1537
(13) Matyjaszewski, K; Science 1996, V272, P866
(14) Muller, A; Macromolecules 1997, V30, P7015
(15) Simon, P; Polym Prepr (Am Chem Soc, Div Polym Chem) 1997, V38(1), P498
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HCAPLUS

(16) Weimer, M; J Polym Sci Polym Chem 1998, V36, P955 HCAPLUS

(17) Yan, D; Macromolecules 1997, V30, P7024 HCAPLUS

IT 213453-03-3P 213453-06-6P

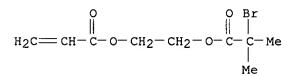
RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (hyperbranched; prepn. of hyperbranched polyacrylates by atom-transfer radical polymn. by using zero-valent copper catalyst)

RN 213453-03-3 HCAPLUS

CN 2-Propenoic acid, 2-(2-bromo-2-methyl-1-oxopropoxy)ethyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 213453-02-2 CMF C9 H13 Br O4



elected species
Claim 2
282

RN 213453-06-6 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 2-(2-bromo-1-oxopropoxy)ethyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 213453-05-5 CMF C9 H13 Br O4

L57 ANSWER 6 OF 6 HCAPLUS COPYRIGHT 2003 ACS

//AN /1998:621247 HCAPLUS

(DN (129:261036

TI Controlled atom or group-transfer radical polymerization, coupling of molecules, multifunctional polymerization initiators, and formation of telechelic functional material

IN Matyjaszewski, Krzysztof; Gaynor, Scott G.; Coca, Simion

PA Carnegie Mellon University, USA

SO / PCT_Int. Appl., 230 pp.

CODEN: PIXXD2

DT Patent

LA English

IC ICM C08F004-10

ICS C08F008-00; C08F008-38

CC 35-3 (Chemistry of Synthetic High Polymers)

FAN.CNT 6

PATENT NO. KIND DATE APPLICATION NO. DATE

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19980917
                                           WO 1998-US4333
                                                           19980311
PΙ
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             KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX,
             NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT,
             UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI,
             FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM,
             GA, GN, ML, MR, NE, SN, TD, TG
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PRAI US 1997-39543P
                       Ρ
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                       Р
                            19970402
     US 1998-18554
                      Α
                            19980204
     US 1995-559309
                       Α3
                            19951115
     WO 1998-US4333
                      W
                            19980311
     A process for ATRP polymn. and coupling of mols. by radical processes is
AΒ
     improved by the selection of various ligands, counterions, transition
     metal compds. and/or zero oxidn. state transition metals to give improved
     control over mol. wt., mol. wt. distribution, functionality and compns. of
     the products formed. The process is useful not only in polymn. but also
     in coupling of mols. of any size, by generation and coupling of the
     appropriate radicals, and in modifying chain ends of functionalized
     polymers. Thus, styrene was bulk polymd. in the presence of iron powder,
     DMF, and 1-(bromoethyl)benzene for 9 h at 110.degree. with 85% conversion
     to polymer having Mn 8960 and Mw/Mn 1.33.
ST
     group transfer radical polymn vinyl compd; catalyst radical polymn
     coupling mol; polystyrene prepn catalyst; bromoethylbenzene initiator
     group transfer radical polymn; iron catalyst group transfer radical
     polymn; DMF ligand group transfer radical polymn
IT
     Alcohols, preparation
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (amino, prepn. of; controlled atom or group-transfer radical polymn.,
        coupling of mols., multifunctional polymn. initiators, and formation of
        telechelic functional material)
TT
     Ligands
     RL: CAT (Catalyst use); USES (Uses)
        (controlled atom or group-transfer radical polymn., coupling of mols.,
        multifunctional polymn. initiators, and formation of telechelic
        functional material)
IT
     Polymerization
     Polymerization catalysts
        (group-transfer; controlled atom or group-transfer radical polymn.,
        coupling of mols., multifunctional polymn. initiators, and formation of
        telechelic functional material)
ΙT
     Polymers, preparation
     RL: IMF (Industrial manufacture); PREP (Preparation)
        (hyperbranched or bottle-brush, prepn. of; controlled atom or
        group-transfer radical polymn., coupling of mols., multifunctional
        polymn. initiators, and formation of telechelic functional material)
IT
     Silsesquioxanes
     RL: CAT (Catalyst use); USES (Uses)
        (initiator; controlled atom or group-transfer radical polymn., coupling
        of mols., multifunctional polymn. initiators, and formation of
```

telechelic functional material)

Polysiloxanes, preparation IT RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses) (macroinitiators and macromonomers; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) IT Amines, preparation RL: SPN (Synthetic preparation); PREP (Preparation) (primary, vinyl polymers terminated with; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) IT Amines, preparation RL: SPN (Synthetic preparation); PREP (Preparation) (secondary, vinyl polymers terminated with; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) IT Polymers, preparation RL: IMF (Industrial manufacture); PREP (Preparation) (star-branched, prepn. from bromine-terminated polystyrene and divinylbenzene; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) TΨ Macromonomers RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (terminated siloxanes; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) ΙT Crosslinking (thermal; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) IT 7631-86-9, Silica, uses RL: CAT (Catalyst use); USES (Uses) (activated with silanes, catalyst support; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) ΙT 78-67-1, AIBN 1313-13-9, Manganese oxide, uses 7439-95-4, Magnesium, 7439-96-5, Manganese, uses 7440-02-0, Nickel, uses Ruthenium, uses 7440-19-9, Samarium, uses 7440-2, 7440-47-3, Chromium, uses 7440-50-8, Copper, uses 7440-22-4, Silver, uses 7447-39-4, Copper (II) chloride, uses 7773-01-5, Manganese chloride 7787-70-4, Copper (I) bromide 7789-45-9, Copper (II) bromide 7789-46-0, Iron (II) 10025-73-7, Chromium trichloride 10031-26-2, Iron tribromide 10049-05-5, Chromium dichloride 12597-70-5, Copper bronze 26490-65-3, Cuprous hexafluorophosphate 34946-82-2, Copper ditriflate 37234-97-2 126949-65-3 RL: CAT (Catalyst use); USES (Uses) (catalyst; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) 5468-93-9DP, polystyrene terminated with ΙT 85673-60-5DP, polystyrene terminated with RL: IMF (Industrial manufacture); PREP (Preparation) (controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material)

- PASTERCZYK 09/359359 Page 23 4648-54-8D, Trimethylsilyl azide, polystyrene terminated with IT RL: NUU (Other use, unclassified); USES (Uses) (controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) 623-24-5DP, .alpha.,.alpha.'-Dibromo-p-xylene, polystyrene terminated with IT RL: SPN (Synthetic preparation); PREP (Preparation) (controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) IT 2564-83-2, TEMPO RL: RCT (Reactant); RACT (Reactant or reagent) (coupling with alkyl halides; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) 585-71-7, 1-Phenylethyl bromide 600-00-0, Ethyl 2-bromoisobutyrate 5445-17-0, Methyl 2-bromopropionate 19481-82-4, 2-Bromopropionitrile RL: CAT (Catalyst use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses) (initiator and coupling with tetramethylpiperidinyloxy; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) ΙT 213453-21-5P

 - RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(initiator, prepn. of; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material)

IT 79-07-2, 2-Chloroacetamide 80-58-0, 2-Bromobutyric acid 98-59-9, Tosyl 104-81-4, 4-Methylbenzyl bromide 106-95-6, Allyl bromide, chloride 124-63-0, Methanesulfonyl chloride 535-11-5, Ethyl 2-bromopropionate 589-15-1, 4-Bromobenzyl bromide 590-17-0, 598-54-9, Copper monoacetate 672-65-1, 1-Phenylethyl Bromoacetonitrile 776-74-9, Bromodiphenylmethane 1643-19-2, Tetrabutylammonium chloride 2052-01-9, 2-Bromoisobutyric acid 2549-51-1, Vinyl bromide 2916-14-5, Allyl chloroacetate 3012-37-1, Benzyl chloroacetate 3042-81-7, Methyl .alpha.-bromophenyl acetate thiocvanate 5061-21-2 17201-43-3, 4-Cyanobenzyl bromide 17639-93-9, Methyl 5468-93-9 18301-66-1 29263-94-3, Diethyl 2-bromo-2-2-chloropropionate methylmalonate 39149-80-9, tert-Butyl 2-bromopropionate 56905-18-1, Methyl 2-iodopropionate 68986-76-5, Copper (I) 2-thiophenecarboxylate 82280-42-0 87129-38-2D, Allyl-2-bromopropionate, reaction products with cyclosiloxanes 208446-93-9 213137-90-7 213453-16-8D, reaction products with allylbromopropionate RL: CAT (Catalyst use); USES (Uses)

(initiator; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material)

- IT 603-35-0, Triphenyl phosphine, uses 102-82-9, Tributyl amine RL: CAT (Catalyst use); USES (Uses)
 - (ligand, catalyst; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material)
- IT 17678-99-8P 33527-91-2P 204580-80-3P RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)
 - (ligand, prepn. of; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of

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telechelic functional material)
     68-12-2, DMF, uses 110-18-9 148-24-3, 8-Hydroxyquinoline, uses
ΙT
                                1116-76-3, Tri(n-octylamine)
     366-18-7, 2,2'-Bipyridine
                                                                2085-33-8
     2212-32-0, 2-{[2-(Dimethylamino)ethyl]methylamino}ethanol
                                                                 3030-47-5,
     N, N, N', N', N''-Pentamethyldiethylenetriamine 3083-10-1,
     1,1,4,7,10,10-Hexamethyltriethylenetetramine
                                                   41203-22-9,
     1,4,8,11-Tetramethyl-1,4,8,11-tetraazacyclotetradecane 72230-93-4,
     4,4'-Di(5-nonyl)-2,2'-bipyridine
                                        72914-19-3
     RL: CAT (Catalyst use); USES (Uses)
        (ligand; controlled atom or group-transfer radical polymn., coupling of
        mols., multifunctional polymn. initiators, and formation of telechelic
        functional material)
IT
     116629-00-6P
     RL: CAT (Catalyst use); RCT (Reactant); SPN (Synthetic preparation); PREP
     (Preparation); RACT (Reactant or reagent); USES (Uses)
        (macroinitiator, prepn. and reaction of; controlled atom or
        group-transfer radical polymn., coupling of mols., multifunctional
        polymn. initiators, and formation of telechelic functional material)
ΙT
     213453-20-4P, 2,2'-Azobis[2-methyl-N-(2-(2-bromoisobutyryloxy)ethyl)propio
     namide-styrene copolymer
     RL: CAT (Catalyst use); IMF (Industrial manufacture); PREP (Preparation);
     USES (Uses)
        (macroinitiator, prepn. of; controlled atom or group-transfer radical
        polymn., coupling of mols., multifunctional polymn. initiators, and
        formation of telechelic functional material)
TT
     74143-32-1DP, poly(hexamethylcyclotrisiloxane) terminated with
     99349-00-5DP, poly(hexamethylcyclotrisiloxane) terminated with
     213453-17-9P, Ally1-2-bromopropionate-2,4,6,8-tetramethylcyclosiloxane
     copolymer
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
        (macroinitiator, prepn. of; controlled atom or group-transfer radical
        polymn., coupling of mols., multifunctional polymn. initiators, and
        formation of telechelic functional material)
IT
     9003-27-4DP, Polyisobutene, styrene-terminated
                                                      25038-76-0P.
     Polynorbornene 25038-78-2P, Dicyclopentadiene homopolymer
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (macroinitiator, prepn. of; controlled atom or group-transfer radical
        polymn., coupling of mols., multifunctional polymn. initiators, and
        formation of telechelic functional material)
IT
     25084-99-5DP, Hexamethylcyclotrisiloxane homopolymer,
     (chlorodimethylsilylethyl)styrene- or [(chloromethyl)phenylethyl]dimethylc
     hlorosilane-terminated
     RL: CAT (Catalyst use); RCT (Reactant); SPN (Synthetic preparation); PREP
     (Preparation); RACT (Reactant or reagent); USES (Uses)
        (macromonomer or macroinitiator; controlled atom or group-transfer
        radical polymn., coupling of mols., multifunctional polymn. initiators,
        and formation of telechelic functional material)
ΙT
     5958-97-4DP, poly(hexamethylcyclotrisiloxane) terminated with
     213453-18-ODP, poly(hexamethylcyclotrisiloxane) terminated with
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (macromonomer; controlled atom or group-transfer radical polymn.,
        coupling of mols., multifunctional polymn. initiators, and formation of
        telechelic functional material)
IT
     71-43-2, Benzene, miscellaneous
                                     96-49-1, Ethylene carbonate
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     108-32-7
     RL: MSC (Miscellaneous)
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(polymn. solvent; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) 7439-89-6, Iron, uses RL: CAT (Catalyst use); USES (Uses) (powder, catalyst; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) 9003-53-6DP, Polystyrene, bromine-terminated RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent) (prepn. and coupling of; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) 213453-19-1P RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent) (prepn. and polymn. of; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) 9003-21-8, Methyl acrylate homopolymer RL: CAT (Catalyst use); USES (Uses) (prepn. of; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material) 80-62-6DP, graft copolymer with chlorosulfonated polyethylene 100-42-5DP, graft copolymer with chlorosulfonated polyethylene 109-73-9DP, n-Butylamine, vinyl polymers terminated with Isobornyl acrylate, graft copolymers with brominated rubbers 9002-88-4DP, chlorosulfonated, graft polymer with vinyl compds. 9003-21-8DP, Methyl acrylate homopolymer, functional group-terminated 9003-49-0P, Butyl acrylate homopolymer 9003-53-6P, Polystyrene 9003-54-7P, Acrylonitrile-styrene copolymer 9010-85-9DP, Isobutene-isoprene copolymer, brominated, graft polymers with vinyl 9011-14-7P, Poly(methyl methacrylate) 13325-10-5DP, 4-Aminobutanol, vinyl polymers terminated with 25014-41-9P, Acrylonitrile homopolymer 25067-63-4P, Methyl acrylate-vinyl acetate copolymer 25154-86-3P, 2-(Dimethylamino)ethyl methacrylate homopolymer 25213-17-6P, Acrylonitrile-isobutene copolymer 25232-27-3P, tert-Butyl acrylate homopolymer 25249-16-5P, 2-Hydroxyethyl methacrylate homopolymer 25266-62-0P, Allyl acrylate polymer 25767-47-9P, Butyl acrylate-styrene copolymer 26022-14-0P, 2-Hydroxyethyl acrylate polymer 26374-91-4P, Glycidyl acrylate homopolymer 26588-80-7P, Butyl acrylate-2-hydroxyethyl methacrylate-methyl methacrylate-styrene copolymer 26628-22-8DP, Sodium azide, vinyl polymers terminated with 30323-87-6P, Isobornyl acrylate 30811-69-9P, Vinyl acrylate polymer homopolymer 31049-58-8P, Butyl acrylate-isobutylene copolymer 40704-75-4P, N-(2-Hydroxypropyl) methacrylamide polymer 56467-21-1P, Butyl acrylate-3-(trimethoxysilyl)propyl methacrylate copolymer 61128-14-1DP, Isobutylene-p-methylstyrene copolymer, brominated, graft copolymers with vinyl compds. 107227-34-9P, Acrylonitrile-isobutylene alternating copolymer 108150-11-4P, Methyl acrylate-methyl methacrylate block copolymer 108501-18-4P, Butyl acrylate-methyl methacrylate block copolymer 110772-34-4P, Butyl acrylate-styrene block copolymer 112965-31-8P, Acrylonitrile-butyl acrylate-styrene block copolymer

121264-61-7P, Butyl acrylate-2-hydroxyethyl acrylate block copolymer 136234-79-2P, N-Cyclohexyl maleimide-styrene alternating copolymer

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137168-27-5P, Acrylonitrile-butyl acrylate block copolymer
                                                                  144719-01-7P,
     Methyl acrylate-methyl methacrylate-styrene block copolymer
     166664-50-2P, Butyl acrylate-isobutylene alternating copolymer
     188065-74-9P, 2-(2-Bromopropionyloxy)ethyl acrylate homopolymer
                                 213453-09-9P
                                                213453-12-4P,
     213453-03-3P 213453-06-6P
     Styrene-vinyl chloride-vinyl chloroacetate graft copolymer
                                                                  213453-13-5P,
     Butyl acrylate-2-ethylhexyl acrylate-Zonyl TAN block copolymer
                    213555-59-0P, Acrylonitrile-methyl acrylate block copolymer
     RL: IMF (Industrial manufacture); PREP (Preparation)
        (prepn. of; controlled atom or group-transfer radical polymn., coupling
        of mols., multifunctional polymn. initiators, and formation of
        telechelic functional material)
                    212128-87-5P
                                   212128-91-1P
                                                  212129-00-5P
     154554-67-3P
                                                                 213453-14-6P,
     Isobornyl acrylate-isobutene-methyl acrylate-methyl methacrylate-styrene
     block copolymer
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (prepn. of; controlled atom or group-transfer radical polymn., coupling
        of mols., multifunctional polymn. initiators, and formation of
        telechelic functional material)
     107-21-1, 1,2-Ethanediol, reactions
                                           563-76-8
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reactant in initiator prepn.; controlled atom or group-transfer
        radical polymn., coupling of mols., multifunctional polymn. initiators,
        and formation of telechelic functional material)
     49864-98-4, Hexakis (4-hydroxymethylphenoxy) cyclotriphosphazene
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reactant; controlled atom or group-transfer radical polymn., coupling
        of mols., multifunctional polymn. initiators, and formation of
        telechelic functional material)
     75-09-2, reactions
                          20769-85-1, 2-Bromoisobutyryl bromide
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reaction with azobis[methyl(hydroxyethyl)propionamide]; controlled
        atom or group-transfer radical polymn., coupling of mols.,
        multifunctional polymn. initiators, and formation of telechelic
        functional material)
     61551-69-7, 2,2'-Azobis[2-methyl-N-(2-hydroxyethyl)propionamide
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reaction with haloalkyl halide; controlled atom or group-transfer
        radical polymn., coupling of mols., multifunctional polymn. initiators,
        and formation of telechelic functional material)
     50975-76-3
                  103526-27-8
     RL: NUU (Other use, unclassified); USES (Uses)
        (silica activated by; controlled atom or group-transfer radical
        polymn., coupling of mols., multifunctional polymn. initiators, and
        formation of telechelic functional material)
     100-66-3, Anisole, miscellaneous
                                       25321-22-6, Dichlorobenzene
     RL: MSC (Miscellaneous)
        (solvent; controlled atom or group-transfer radical polymn., coupling
        of mols., multifunctional polymn. initiators, and formation of
        telechelic functional material)
RE.CNT
             THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
(1) Decker; US 3862978 A 1975 HCAPLUS
(2) Patten; Science 1996, V272, P866 HCAPLUS
     213453-21-5P
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
        (initiator, prepn. of; controlled atom or group-transfer radical
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polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material)

RN 213453-21-5 HCAPLUS

CN Benzoic acid, 4-(chloromethyl)-, azobis[(2,2-dimethyl-1-oxo-2,1-ethanediyl)imino-2,1-ethanediyl] ester (9CI) (CA INDEX NAME)

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AMCBP

PAGE 1-B

IT 25067-63-4P, Methyl acrylate-vinyl acetate copolymer 213453-03-3P 213453-06-6P

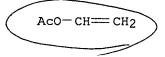
RL: IMF (Industrial manufacture); PREP (Preparation) (prepn. of; controlled atom or group-transfer radical polymn., coupling of mols., multifunctional polymn. initiators, and formation of telechelic functional material)

RN 25067-63-4 HCAPLUS

CN 2-Propenoic acid, methyl ester, polymer with ethenyl acetate (9CI) (CA INDEX NAME)

CM 1

CRN 108-05-4 CMF C4 H6 O2



CM 2

CRN 96-33-3 CMF C4 H6 O2

RN 213453-03-3 HCAPLUS

CN 2-Propenoic acid, 2-(2-bromo-2-methyl-1-oxopropoxy)ethyl ester,

KATHLEEN FULLER EIC 1700/PARKER LAW 308-4290

homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 213453-02-2 CMF C9 H13 Br O4 elected peries #282

RN 213453-06-6 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 2-(2-bromo-1-oxopropoxy)ethyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 213453-05-5 CMF C9 H13 Br O4

=> FILE REG

FILE 'REGISTRY' ENTERED AT 16:27:05 ON 13 MAY 2003 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

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Experimental and calculated property data are now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details: http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf

=> L36;D L44;D L45;D L49; D L53;D L54;D L55 L36 IS NOT A RECOGNIZED COMMAND

COMMAND STACK INTERRUPTED. ENTER "DISPLAY HISTORY" TO SEE WHICH COMMANDS WERE EXECUTED.

The previous command name entered was not recognized by the system. For a list of commands available to you in the current file, enter "HELP COMMANDS" at an arrow prompt (=>).

=> D L36; D L44; D L45; D L49; D L53; D L54; D L55

L36 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2003 ACS

RN 213453-21-5 REGISTRY

CN Benzoic acid, 4-(chloromethyl)-, azobis[(2,2-dimethyl-1-oxo-2,1-ethanediyl)imino-2,1-ethanediyl] ester (9CI) (CA INDEX NAME)
OTHER NAMES:

CN 2,2'-Azobis[2-methyl-N-[2-[4-(chloromethyl)benzoyloxy]ethyl]propionam ide]

FS 3D CONCORD

MF C28 H34 C12 N4 O6

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

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PAGE 1-B

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

4 REFERENCES IN FILE CA (1957 TO DATE)
4 REFERENCES IN FILE CAPLUS (1957 TO DATE)

L44 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2003 ACS

RN 371771-79-8 REGISTRY

CN Butanoic acid, 2-bromo-, 2-[(1-oxo-2-propenyl)oxy]ethyl ester, homopolymer (9CI) (CA INDEX NAME)

MF (C9 H13 Br O4)x

CI PMS

PCT Polyacrylic

SR CA

LC STN Files: CA, CAPLUS

CM 1

CRN 371771-77-6 CMF C9 H13 Br O4

1 REFERENCES IN FILE CA (1957 TO DATE)

1 REFERENCES IN FILE CAPLUS (1957 TO DATE)

L45 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2003 ACS

RN 371771-79-8 REGISTRY

CN Butanoic acid, 2-bromo-, 2-[(1-oxo-2-propenyl)oxy]ethyl ester, homopolymer (9CI) (CA INDEX NAME)

MF (C9 H13 Br O4)x

CI PMS

PCT Polyacrylic

SR CA

LC STN Files: CA, CAPLUS

CM 1

CRN 371771-77-6 CMF C9 H13 Br O4

$$\begin{array}{c|cccc} O & O & Br \\ || & || & || \\ H_2C & \hline \\ CH & C-O-CH_2-CH_2-O-C-CH-Et \\ \end{array}$$

1 REFERENCES IN FILE CA (1957 TO DATE)

1 REFERENCES IN FILE CAPLUS (1957 TO DATE)

L49 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2003 ACS

RN **371771-77-6** REGISTRY

CN Butanoic acid, 2-bromo-, 2-[(1-oxo-2-propenyl)oxy]ethyl ester (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C9 H13 Br O4

CI COM

SR CA

LC STN Files: CA, CAPLUS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1957 TO DATE)

1 REFERENCES IN FILE CAPLUS (1957 TO DATE)

L53 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2003 ACS

RN 213453-03-3 REGISTRY

CN 2-Propenoic acid, 2-(2-bromo-2-methyl-1-oxopropoxy)ethyl ester, homopolymer (9CI) (CA INDEX NAME)

MF (C9 H13 Br O4)x

CI PMS

PCT Polyacrylic

SR CA

LC STN Files: CA, CAPLUS

CM 1

CRN 213453-02-2 CMF C9 H13 Br O4

2 REFERENCES IN FILE CA (1957 TO DATE)

2 REFERENCES IN FILE CAPLUS (1957 TO DATE)

L54 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2003 ACS

RN 213453-03-3 REGISTRY

CN 2-Propenoic acid, 2-(2-bromo-2-methyl-1-oxopropoxy)ethyl ester, homopolymer (9CI) (CA INDEX NAME)

MF (C9 H13 Br O4)x

CI PMS

PCT Polyacrylic

SR CA

LC STN Files: CA, CAPLUS

CM 1

CRN 213453-02-2 CMF C9 H13 Br O4

2 REFERENCES IN FILE CA (1957 TO DATE)

2 REFERENCES IN FILE CAPLUS (1957 TO DATE)

L55 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2003 ACS

RN **213453-02-2** REGISTRY

CN 2-Propenoic acid, 2-(2-bromo-2-methyl-1-oxopropoxy)ethyl ester (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C9 H13 Br O4

CI COM

SR CA

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PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT